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NEWS 4 FEB 02	GENBANK enhanced with SET PLURALS and SET SPELLING
NEWS 5 FEB 06	Patent sequence location (PSL) data added to USGENE
NEWS 6 FEB 10	COMPENDEX reloaded and enhanced
NEWS 7 FEB 11	WTEXTILES reloaded and enhanced
NEWS 8 FEB 19	New patent-examiner citations in 300,000 CA/CAplus patent records provide insights into related prior art
NEWS 9 FEB 19	Increase the precision of your patent queries -- use terms from the IPC Thesaurus, Version 2009.01
NEWS 10 FEB 23	Several formats for image display and print options discontinued in USPATFULL and USPAT2
NEWS 11 FEB 23	MEDLINE now offers more precise author group fields and 2009 MeSH terms
NEWS 12 FEB 23	TOXCENTER updates mirror those of MEDLINE - more precise author group fields and 2009 MeSH terms
NEWS 13 FEB 23	Three million new patent records blast AEROSPACE into STN patent clusters
NEWS 14 FEB 25	USGENE enhanced with patent family and legal status display data from INPADOCDB
NEWS 15 MAR 06	INPADOCDB and INPAFAMDB enhanced with new display formats
NEWS 16 MAR 11	EPFULL backfile enhanced with additional full-text applications and grants
NEWS 17 MAR 11	ESBIOBASE reloaded and enhanced
NEWS 18 MAR 20	CAS databases on STN enhanced with new super role for nanomaterial substances
NEWS 19 MAR 23	CA/CAplus enhanced with more than 250,000 patent equivalents from China
NEWS 20 MAR 30	IMSPATENTS reloaded and enhanced
NEWS 21 APR 03	CAS coverage of exemplified prophetic substances enhanced
NEWS 22 APR 07	STN is raising the limits on saved answers
NEWS 23 APR 24	CA/CAplus now has more comprehensive patent assignee information
NEWS 24 APR 26	USPATFULL and USPAT2 enhanced with patent assignment/reassignment information
NEWS 25 APR 28	CAS patent authority coverage expanded
NEWS 26 APR 28	ENCOMPLIT/ENCOMPLIT2 search fields enhanced
NEWS 27 APR 28	Limits doubled for structure searching in CAS

## REGISTRY

NEWS 28 MAY 08 STN Express, Version 8.4, now available  
NEWS 29 MAY 11 STN on the Web enhanced  
NEWS 30 MAY 11 BEILSTEIN substance information now available on  
STN Easy

NEWS EXPRESS MAY 08 09 CURRENT WINDOWS VERSION IS V8.4,  
AND CURRENT DISCOVER FILE IS DATED 06 APRIL 2009.

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FILE 'REGISTRY' ENTERED AT 08:38:47 ON 11 MAY 2009  
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STRUCTURE FILE UPDATES: 8 MAY 2009 HIGHEST RN 1144618-76-7  
DICTIONARY FILE UPDATES: 8 MAY 2009 HIGHEST RN 1144618-76-7

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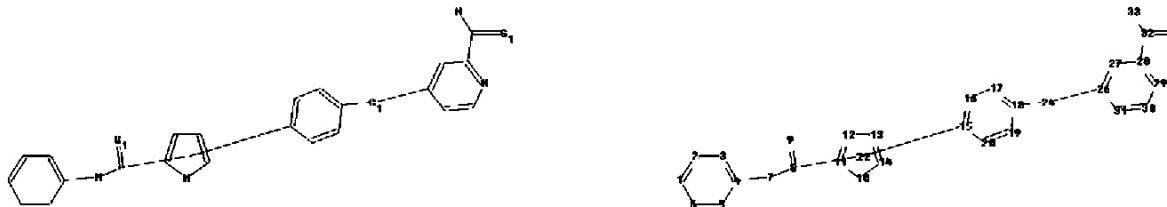
TSCA INFORMATION NOW CURRENT THROUGH January 9, 2009.

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<http://www.cas.org/support/stn/gen/stndoc/properties.html>

=> Uploading C:\Program Files\STNEXP\Queries\10579825 updated.str



chain nodes :

7 8 9 24 32 33 34

ring nodes :

1 2 3 4 5 6 10 11 12 13 14 15 16 17 18 19 20 26 27 28 29 30  
31

chain bonds :

4-7 7-8 8-9 18-24 24-26 28-32 32-33 32-34

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 10-11 10-14 11-12 12-13 13-14 15-16 15-20 16-  
17  
17-18 18-19 19-20 26-27 26-31 27-28 28-29 29-30 30-31

exact/norm bonds :

4-7 7-8 8-9 10-11 10-14 11-12 12-13 13-14 18-24 24-26 32-33 32-34

exact bonds :

28-32

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 15-16 15-20 16-17 17-18 18-19 19-20 26-27 26-  
31

27-28 28-29 29-30 30-31

isolated ring systems :

containing 10 : 26 :

G1:0,S

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:Atom  
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom  
20:Atom 21:Atom  
22:Atom 24:CLASS 26:Atom 27:Atom 28:Atom 29:Atom 30:Atom 31:Atom 32:CLASS  
33:CLASS 34:CLASS

L1 STRUCTURE UPLOADED

=> d L1

L1 HAS NO ANSWERS

L1 STR

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

Structure attributes must be viewed using STN Express query preparation.

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=> file caplus
COST IN U.S. DOLLARS          SINCE FILE      TOTAL
                                ENTRY        SESSION
FULL ESTIMATED COST          0.48          0.70
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FILE 'CAPLUS' ENTERED AT 08:39:06 ON 11 MAY 2009  
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FILE COVERS 1907 - 11 May 2009 VOL 150 ISS 20  
FILE LAST UPDATED: 8 May 2009 (20090508/ED)  
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2009  
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2009

CAplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

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This file contains CAS Registry Numbers for easy and accurate

```
=> s L1 SSS full
REG1stRY INITIATED
Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.
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FULL SEARCH INITIATED 08:39:11 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 3229 TO ITERATE

100.0% PROCESSED 3229 ITERATIONS 5 ANSWERS
SEARCH TIME: 00.00.01

L2 5 SEA SSS FUL L1

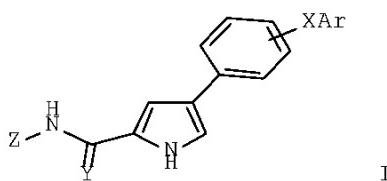
L3 1 L2

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=> d ibib abs hitstr 1-
YOU HAVE REQUESTED DATA FROM 1 ANSWERS - CONTINUE? Y/(N):y
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L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2009 ACS on STN  
 ACCESSION NUMBER: 2005:469894 CAPLUS Full-text  
 DOCUMENT NUMBER: 143:7592  
 TITLE: Preparation of arylpyrrolecarboxamides as Raf kinase inhibitors for treatment of tumors.  
 INVENTOR(S): Finsinger, Dirk; Buchstaller, Hans-Peter; Burgdorf, Lars; Wiesner, Matthias; Amendt, Christiane; Grell, Matthias; Sirrenberg, Christian; Zenke, Frank  
 PATENT ASSIGNEE(S): Merck Patent G.m.b.H., Germany  
 SOURCE: Ger. Offen., 32 pp.  
 CODEN: GWXXBX  
 DOCUMENT TYPE: Patent  
 LANGUAGE: German  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10354060	A1	20050602	DE 2003-10354060	20031119
AU 2004291255	A1	20050602	AU 2004-291255	20041026
CA 2546334	A1	20050602	CA 2004-2546334	20041026
WO 2005049603	A1	20050602	WO 2004-EP12076	20041026
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1685125	A1	20060802	EP 2004-790859	20041026
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
CN 1882571	A	20061220	CN 2004-80034345	20041026
BR 2004016690	A	20070130	BR 2004-16690	20041026
JP 2007511553	T	20070510	JP 2006-540216	20041026
IN 2006KN00936	A	20070420	IN 2006-KN936	20060417
MX 2006005478	A	20060811	MX 2006-5478	20060515
KR 2006118492	A	20061123	KR 2006-709552	20060517
US 20070149594	A1	20070628	US 2006-579825	20060517
PRIORITY APPLN. INFO.:			DE 2003-10354060	A 20031119
			WO 2004-EP12076	W 20041026

OTHER SOURCE(S): MARPAT 143:7592  
 GI



AB Title compds. [I; Ar = (substituted) Ph, naphthyl, biphenyl, heterocyclyl; X = O, S, (CH<sub>2</sub>)<sub>n</sub>, CO, (CH<sub>2</sub>)<sub>n</sub>O, (CH<sub>2</sub>)<sub>n</sub>NH, etc.; n = 1-3; Y = O, S, CHNO<sub>2</sub>, C(CN)<sub>2</sub>, NR<sub>4</sub>; R<sub>4</sub> = H, cyano, OH, etc.; Z = Ar, ArXAr, CH<sub>2</sub>Ar, CH<sub>2</sub>ArXAr; Ar = (substituted) Ph], were prepared as Raf kinase inhibitors (no data). Thus, 4-(PhCH<sub>2</sub>O)C<sub>6</sub>H<sub>4</sub>CH<sub>2</sub>CO<sub>2</sub>H, DMF, and POCl<sub>3</sub> were heated together at 70° for 4 h followed by cooling and addition of ice water and aqueous NaClO<sub>4</sub> to give 98% [2-(4-benzyloxyphenyl)-3-dimethylaminoallylidene]dimethylammonium perchlorate. This was refluxed 24 h with glycine Et ester hydrochloride in EtOH containing 20% NaOEt to give 91% Et 4-(4-benzyloxyphenyl)-1H-pyrrole-2-carboxylate. Hydrogenolysis of the latter in EtOAc over Pd/C gave 91% Et 4-(4-hydroxyphenyl)-1H-pyrrole-2-carboxylate. This was heated with 4-chloropyridine-2-carboxylic acid N-methylamide at 160° for 48 h to give 40% Et 4-[4-(2-methylcarbamoylpyridin-4-yloxy)phenyl]-1H-pyrrole-2-carboxylate. Saponification with 2N NaOH in EtOH at 60° for 16 h followed by acidification with HCl gave 85% free acid, which was stirred 48 h in DMF with 5-amino-2-chlorobenzotrifluoride, N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride, and 1-hydroxybenzotriazole hydrate to give 17% 4-[4-[5-(4-chloro-3-trifluoromethylphenylcarbamoyl)-1H-pyrrol-3-yl]phenoxy]pyridine-2-carboxylic acid N-methylamide.

IT 852455-19-7P 852455-21-1P 852455-22-2P

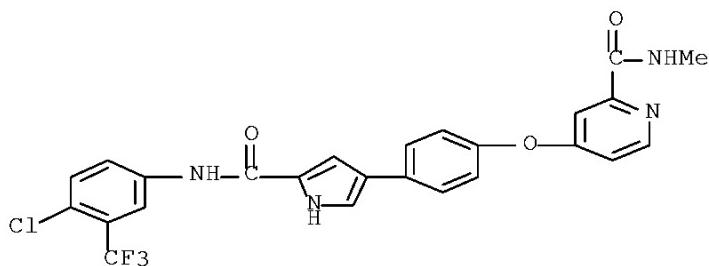
852455-24-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(claimed compound; preparation of arylpyrrolecarboxamides as Raf kinase inhibitors for treatment of tumors)

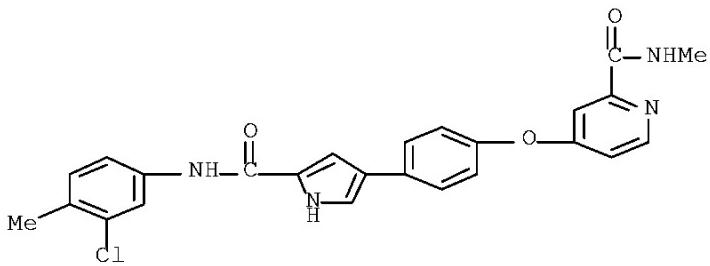
RN 852455-19-7 CAPLUS

CN 2-Pyridinecarboxamide, 4-[4-[5-[[[4-chloro-3-(trifluoromethyl)phenyl]amino]carbonyl]-1H-pyrrol-3-yl]phenoxy]-N-methyl- (CA INDEX NAME)



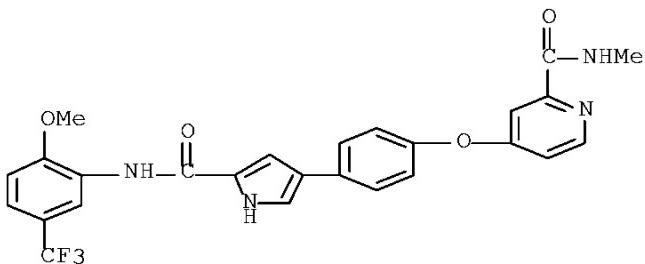
RN 852455-21-1 CAPLUS

CN 2-Pyridinecarboxamide, 4-[4-[5-[(3-chloro-4-methylphenyl)amino]carbonyl]-1H-pyrrol-3-yl]phenoxy]-N-methyl- (CA INDEX NAME)



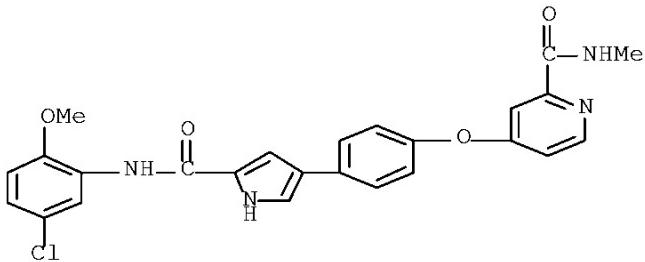
RN 852455-22-2 CAPLUS

CN 2-Pyridinecarboxamide, 4-[4-[5-[[[2-methoxy-5-(trifluoromethyl)phenyl]amino]carbonyl]-1H-pyrrol-3-yl]phenoxy]-N-methyl-  
(CA INDEX NAME)



RN 852455-24-4 CAPLUS

CN 2-Pyridinecarboxamide, 4-[4-[5-[[[5-chloro-2-methoxyphenyl]amino]carbonyl]-1H-pyrrol-3-yl]phenoxy]-N-methyl-  
(CA INDEX NAME)



=> log off

ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y)/N/HOLD:y

STN INTERNATIONAL LOGOFF AT 08:40:04 ON 11 MAY 2009